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APPLICATION NO.	FILIT	NG DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.	
10/699,110	10/	30/2003	Jerome B. Zeldis	9516-083-999	9516-083-999 1866	
20583 JONES DAY	7590	05/04/2007		EXAMINER		
222 EAST 41ST ST				FAY, ZOHREH A		
NEW YORK,	NY 10017			ART UNIT PAPER NUMBER		
				1618		
				MAIL DATE	DELIVERY MODE	
				05/04/2007	PAPER	

Please find below and/or attached an Office communication concerning this application or proceeding.

The time period for reply, if any, is set in the attached communication.

		Application No.	Applicant(s)
Office Action Summary		10/699,110	ZELDIS, JEROME B.
		Examiner	Art Unit
		Zohreh A. Fay	1618
The N Period for Reply	IAILING DATE of this communication app	ears on the cover sheet with the c	orrespondence address
A SHORTEN WHICHEVEF - Extensions of ti after SIX (6) MC - If NO period for - Failure to reply Any reply receive	IED STATUTORY PERIOD FOR REPLY R IS LONGER, FROM THE MAILING DA me may be available under the provisions of 37 CFR 1.13 DNTHS from the mailing date of this communication. reply is specified above, the maximum statutory period within the set or extended period for reply will, by statute, red by the Office later than three months after the mailing erm adjustment. See 37 CFR 1.704(b).	ATE OF THIS COMMUNICATION 36(a). In no event, however, may a reply be tim vill apply and will expire SIX (6) MONTHS from cause the application to become ABANDONEI	Lely filed the mailing date of this communication. D (35 U.S.C. § 133).
Status			
2a) ☐ This ac 3) ☐ Since t	nsive to communication(s) filed on ction is FINAL . 2b)⊠ This this application is in condition for allowar in accordance with the practice under <i>E</i>	action is non-final. nce except for formal matters, pro	
Disposition of C	Claims		
4a) Of t 5) ☐ Claim(6) ☑ Claim(7) ☐ Claim(s) <u>1-7,18 and 19</u> is/are pending in the apthe above claim(s) is/are withdraves) is/are allowed. s) <u>1-7,18 and 19</u> is/are rejected. s) is/are objected to. s) are subject to restriction and/or	vn from consideration.	
Application Pap	ers		•
10)☐ The dra Applica Replace	ecification is objected to by the Examiner twing(s) filed on is/are: a) accept and a section to the common drawing sheet(s) including the correction or declaration is objected to by the Examiner.	epted or b) objected to by the Edrawing(s) be held in abeyance. See ion is required if the drawing(s) is obj	e 37 CFR 1.85(a). ected to. See 37 CFR 1.121(d).
Priority under 3	5 U.S.C. § 119		
a)	viedgment is made of a claim for foreign b) Some * c) None of: Certified copies of the priority documents Certified copies of the priority documents Copies of the certified copies of the priority Application from the International Bureau attached detailed Office action for a list of	s have been received. s have been received in Application ity documents have been received (PCT Rule 17.2(a)).	on No ed in this National Stage
Attachment(s)			
1) Notice of Refe 2) Notice of Draff	rences Cited (PTO-892) sperson's Patent Drawing Review (PTO-948) sclosure Statement(s) (PTO/SB/08) lail Date	4) Interview Summary Paper No(s)/Mail Da 5) Notice of Informal P 6) Other:	ite

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Claims 1-7 and 18-19 are presented for examination.

The response to the restriction of requirement of January 18, 2007 has been received and entered.

The following is a quotation of the first paragraph of 35 U.S.C. 112:

The specification shall contain a written description of the invention, and of the manner and process of making and using it, in such full, clear, concise, and exact terms as to enable any person skilled in the art to which it pertains, or with which it is most nearly connected, to make and use the same and shall set forth the best mode contemplated by the inventor of carrying out his invention.

Claims 1-7 and 18-19 are rejected under 35 U.S.C. 112, first paragraph, because the specification, while being enabling for treating macular degeneration using the claimed compound, does not reasonably provide enablement for "preventing" macular degeneration using the claimed compounds. The specification does not enable any person skilled in the art to which it pertains, or with which it is most nearly connected, to use the invention commensurate in scope with these claims. The factors to be considered whether a disclosure meets the enablement requirements of 35 U.S.C. 112, first paragraph, have been described in In re Wands, 8 USPQ2d 1400 (Fed. Cir.1988). Among these factors are:

1) The nature of the invention:

The claims are drawn to a method of treating macular generation using a selective cytokine inhibitor presented by the compound in claims 1 and 18.

2) The state of the prior art:

The prior art does not recognize that the prevention of macular degeneration is easily accomplished. According to Lance, Current Medical Diagnosis

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and Treatment, 43rd edition, pages 159-160, photodynamic laser therapy is the only known treatment for macular degeneration.

3) The relative skill of those in the art:

The relative skill of those in the art is high.

4) The predictability or unpredictability of the art:

The unpredictability of pharmaceutical and chemical art is high.

5) The breath of the claims:

The claims are very broad and encompass a composition for treating or preventing macular degeneration.

6) The amount of direction or guidance provided:

Applicant's specification provides guidance for and it is only enabled for the treatment of macular degeneration using the claimed compound. In re Dreshfield, 110 F.2d 235, 45 USPQ 36 (CCPA 1940), gives this general rule: "It is well settled that in cases involving chemicals and chemical compounds, which differ radically in their properties it must appear in applicant's specification either by the enumeration of a sufficient number of the members of the group or by other appropriate language, that the chemicals and chemical combinations included in the claims are capable of accomplishing the desired results." Applicant's specification does not set forth any examples to provide support for the prevention of macular degeneration using the claimed compound.

7) The presence or absence of working examples;

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The examples in applicant's specification are drawn to the effect of the

claimed compound on treating macular degeneration.

8) The quantity of experimentation necessary;

Since compound structure and activity for such pharmaceutical use must be determined from case to case by painstaking experimental study, one of ordinary skill in the art would be burdened with undue experimentation to determine the effect of the claimed compound on preventing macular degeneration.

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Zohreh A. Fay whose telephone number is (571) 272-0573. The examiner can normally be reached on Monday to Friday 9:30-6:00.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Michael Hartley can be reached on (571) 272-0616. The fax phone number for the organization where this application or proceeding is assigned is 571-273-8300.

Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see http://pair-direct.uspto.gov. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free). If you would like assistance from a USPTO Customer Service Representative or access to the automated information system, call 800-786-9199 (IN USA OR CANADA) or 571-272-1000.

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